In the Claims:

Please cancel claim 20.

Please amend claims 7, 9-12, 14 and 16-19 as follows:

- 1. (Canceled)
- 2. (Canceled)
- 3. (Canceled)
- 4. (Canceled)
- 5. (Canceled)
- 6. (Canceled)
- 7. (Currently Amended) A method to identify a compound that inhibits wild type full length aggrecanase metalloprotease enzymatic activity comprising the steps of:

comprising a metalloprotease domain, and a peptide substrate, said peptide substrate being less than 40 amino acids in length, wherein the peptide substrate comprises a cleavage site between a glutamic acid on an N-terminal side of the cleavage site and a non-polar or uncharged amino acid residue on a C-terminal side of the cleavage site and wherein the peptide substrate is cleavable by said truncated aggrecanase; and

detecting cleavage of the peptide <u>substrate</u>, wherein inhibition of peptide <u>substrate</u> cleavage in the <u>a</u> presence of a test compound indicates compound inhibition of <u>wild type full</u> <u>length</u> aggrecanase <u>metalloprotease</u> enzymatic activity.

8. (Original) The method of claim 7 wherein the method is conducted in a single reaction vessel.

- 9. (Currently Amended) The method of claim 7 wherein the wild type full length aggrecanase enzyme is selected from the group consisting of aggrecanase-1 and aggrecanase-2 aggrecanase-2.
- 10. (Currently Amended) The method of claim 7 wherein the peptide <u>substrate</u> is selected from the group consisting of SEQ ID NO:3, SEQ ID NO:4, SEQ ID NO:5, SEQ ID NO:6 and SEQ ID NO:7.
- 11. (Currently Amended) The method of claim 7 wherein the peptide <u>substrate</u> further comprises a detectable label selected from the group consisting of ¹²⁵I, ¹³¹I, ³H, ¹⁴C, ³⁵S, ³²P, ³³P, a fluorescent dye, and a colorimetric indicator.
- 12. (Currently Amended) The method of claim 7 wherein the peptide <u>substrate</u> further comprises a fluorophore and a quencher or acceptor located at opposite ends of the cleavage site of the peptide.
- 13. (Previously Amended) The method of claim 7 wherein the truncated aggrecanase is in a cell expressing the truncated aggrecanase.
- 14. (Currently Amended) A method to detect the ability of a compound to inhibit wild type inhibition of metalloprotease enzymatic activity of full length aggrecanase-1 or -2 enzymatic activity comprising the steps of:

cell, said truncated aggrecanase comprising a metalloprotease domain, and a peptide substrate having a detectable label and an amino acid sequence selected from the group consisting of SEQ ID NO:3 and SEQ ID NO:4 in a presence or an absence of a test compound;

incubating the <u>test</u> compound, the truncated aggrecanase and <u>the</u> peptide <u>substrate</u> to <u>permit enzymatic cleavage of the peptide produce a detectable product as a result of metalloprotease enzymatic activity upon the peptide substrate; and</u> measuring a quantity of product formed in a presence of the test compound
and in an absence of the test compound, whereby the inhibition is detected when an amount
of product formed in the presence of the test compound is less than that in the absence of the
test compound.

measuring enzymatic cleavage of the peptide;

wherein said measuring involves determining presence or absence of cleavage of said peptide.

- 15. (Original) The method of claim 14 wherein the peptide comprises a detectable label selected from the group consisting of ¹²⁵I, ¹³¹I, ³H, ¹⁴C, ³⁵S, ³²P, ³³P, a fluorescent dye, or a colorimetric indicator.
- 16. (Currently Amended) The method of claim 14 wherein the peptide <u>substrate</u> comprises a fluorophore and a quencher or acceptor located at opposite ends of the cleavage site of the peptide <u>substrate</u>.
- 17. (Currently Amended) A method to identify a compound capable of inhibiting wild type full length aggrecanase activity comprising the steps;

providing a peptide <u>substrate</u> comprising an affinity moiety, an amino acid sequence selected from a group consisting of SEQ. ID NO:3 and SEQ ID NO:4 and a detectable label, said affinity moiety and label located on opposite sides of a cleavage site encoded by the amino acid sequence;

contacting the peptide <u>substrate</u> with an affinity capture coated solid phase support for sufficient time to bind a portion of the peptide <u>substrate</u>;

washing the support to remove unbound peptide substrate;

contacting a solution comprising a test compound and functional enzyme truncated aggrecanase comprising a metalloprotease domain with the peptide substrate bound solid

phase support for sufficient time to allow enzymatic cleavage of the peptide <u>substrate</u>, thereby releasing the peptide <u>substrate</u> and detectable label into the solution; and

measuring changes in the quantity of the detectable label as a result of compound modulation of expected enzymatic function.

- 18. (Currently Amended) The method of claim 17 wherein the enzyme <u>full length</u> aggrecanase is selected from the group consisting of wild type <u>full length</u> aggrecanase-1 and -2.
- 19. (Currently Amended) The method of claim 17 wherein the peptide <u>substrate</u> comprises a detectable label selected from the group consisting of ¹²⁵I, ¹³¹I, ³H, ¹⁴C, ³⁵S, ³²P, ³³P, a fluorescent dye, or a colorimetric indicator.
 - 20. (Cancelled)
- 21. (Previously Amended) The method of claim 7, wherein said truncated aggrecanase is selected from the group consisting of SEQ ID NO: 8 and SEQ ID NO:9 and homologues thereof.